

Form PTO-1449 U.S. DEPARTMENT OF COMMERCE
(REV. 7-80) PATENT AND TRADEMARK OFFICE

MAR 8 7 2005

U.S. PATENT & TRADEMARK OFFICE

LIST OF PRIOR ART
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(Use several sheets if necessary)

Atty. Docket No. (Optional)

17719 (PC27263A)

Application Number

10/522,254

Applicant(s)

Manuela Villa, et al.

Filing Date

January 25, 2005

Group Art Unit

Unassigned

U.S. PATENT DOCUMENTS

EXAMINER INITIAL*		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE (if appropriate)
	AA						
	AB						

U.S. PATENT APPLICATION PUBLICATIONS

EXAMINER INITIAL*		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE (if appropriate)
DL	AA	2003/0073672 A1	4/17/03	Breitenbucher et al.			
	AB						
	AC						

FOREIGN PATENT DOCUMENTS

	REF	DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION	
							YES	NO
DL		WO 02/064574 A2	8/22/02	PCT				
DL		WO 02/12242 A2	2/14/02	PCT				
DL		WO 96/12720	5/2/96	PCT				

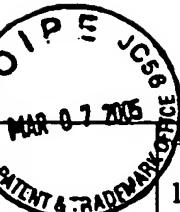
OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

DL	Kikuchi C. et al., "Tetrahydrothienopyridylbutyl-Tetrahydrobenzindoles: New Selective Ligands of the 5-HT Receptor", <i>Bioorganic & Medicinal Chemistry Letters</i> , 12(18):2549-2552 (2002), XP-002256018
DL	Singh P. et al., "Quantitative Structure-Activity Relationship Studies on a New Class of Antihypertensive Agents: Derivatives of 3-Aryl-4,5,6,7-Tetrahydro-1H-Pyrazolo[4,3-c]Pyridine", <i>Quantitative Structure-Activity Relationships</i> , 9(1):29-32 (1990), XP-001155089
DL	Winters G. et al., "Synthesis, in Vitro [³ H]Prazosin Displacement, and in Vivo Activity of 3-Aryl-4,5,6,7-Tetrahydropyrazolo[4,3-c]Pyridines, a New Class of Antihypertensive Agents", <i>J. Med. Chem.</i> , 28(7):934-940 (1985), XP-002256019

EXAMINER		DATE CONSIDERED
		12-04-06

* EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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1 ✓ Radinov R. et al., "3-Phenylpyrazolo[4,3-c]Pyridine and Derivatives: Structure Determination", *Journal of Molecular Structure*, 158:99-108 (1987), XP-009018257

2 ✓ Lackey K. et al., "The Discovery of Potent cRaf1 Kinase Inhibitors", *Bioorganic & Medicinal Chemistry Letters*, 10(3):223-226 (2000), XP-004188821

3 ✓ Philip Cohen, "The Development and Therapeutic Potential of Protein Kinase Inhibitors", *Current Opinion in Chemical Biology*, 3(4):459-465 (1999), XP-002216616

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